

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
27 September 2001 (27.09.2001)

PCT

(10) International Publication Number
WO 01/70206 A2

(51) International Patent Classification⁷: **A61K 31/00**

(21) International Application Number: PCT/HU01/00027

(22) International Filing Date: 1 March 2001 (01.03.2001)

(25) Filing Language: Hungarian

(26) Publication Language: English

(30) Priority Data:
P 00 00997 6 March 2000 (06.03.2000) HU

(71) Applicant and

(72) Inventor: **JÁVOR, András** [HU/HU]; Torbágy utca 8. III. em. 11, H-1118 Budapest (HU).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **PELLIONISZNÉ**

PARÓCZAI, Margit [HU/HU]; Trencsényi utca 45, H-1125 Budapest (HU). **SZÉKELY, Ákosné** [HU/HU]; Wesselényi utca 130, H-1204 Budapest (HU).

(81) Designated States (*national*): HU, JP, US.

(84) Designated States (*regional*): European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.



WO 01/70206 A2

(54) Title: NEW DRUG COMBINATION

(57) Abstract: The invention relates to a drug combination comprising 1:1-30:1 mass ratio blend of lecithin with 10.00-99.99 mass% L-alpha-phosphatidylcholine content and ascorbic acid which is suitable for the preservation and increase of the physical and cognitive performance of healthy and sick people.

NEW DRUG COMBINATION

The present invention relates to the use of a combination of lecithin and ascorbic acid for the preservation or increase of memory and physical performance of healthy and sick people, or for the replacing therapy of the two components, or for the prevention or treatment of all that illnesses in which lipidperoxidation has role.

In the commercial and legal practice the name of lecithin means the mixture of polar and non-polar lipids from vegetable or animal source, containing acetone insoluble components minimum in 60%. About 60% of acetone insoluble part of lecithin is the mixture of phosphor containing lipids, which has as a main component L-alpha-phosphatidylcholine (Drug and Cosmetic Industry, February, 1992.). According to scientific terminology the name of lecithin means only L-alpha-phosphatidylcholine. In the description of the invention the name of lecithin is used in commercial and legal meaning.

In living body inside it in vertebrates and so in human, too, phospholipids and their decomposing products have vital importance role. In human body among phospholipids L-alpha-phosphatidylcholine can be observed in the largest volume, which is the most significant phospholipid from the point of pharmacological and biological view. L-alpha-phosphatidylcholine can be found mainly in membranes of cells, where it is present as a structure component and as an initial compound of different biological active agents. This double role of L-alpha-phosphatidylcholine is essential for the sake of the integrity and functional action of cells (Lecithin and health care, Semmelweis Verlag, Hoya, Germany, 1985; Nutrition Review, Vol. 52, 327-339, 1994.)

It is also important to optimal cell function the chemical structure and ratio of membrane components (proteins, lipids). If the favourable ratio of components changes, for example on the effect of outside factors (malnutrition or unhealthy food intake, physical trauma, bacterial or virus infection) or if the chemical structure of membrane components changes, for example because of peroxidation, they may induce depending on the type of cell and tissue different illnesses, for example vascular wall damage (atherosclerosis).

During the biological ageing the ratio of components of cell membranes and so their structures also change because the increase of cholesterol and decrease of L-alpha-phosphatidylcholine content, which result the decrease of cell function and increase of vulnerability of cells.

One possibility to assure the optimal composition of cell membranes as well as to protect their chemical structure mainly against with peroxidation is that if the main component of membrane phospholipids is present in enough amount and simultaneously
35 membranes are protected from peroxidation with a natural antioxidant such as ascorbic acid.

Lecithin from animal or vegetable source because its complex physiological role has been used by medical science for a long time as a general roborant, at healthy and sick people. Recently, lecithin for this purpose is used alone (Buerlecithin) or in combination with vitamins and trace elements (Gerovit capsule, Kondi tablet Hungarian products). Lecithin,
40 beside its general tonic action is also used for therapeutic aim in some illnesses, for example in ageing memory disturbance caused by acetylcholine deficit (senile dementia) or in moving inco-ordination (tardive dyskinesia). In these cases lecithin promotes as a favourable choline source to the synthesis of acetylcholine. Probably this property of lecithin explains its favourable adjuvant effect in Alzheimer disease (Biological Psychiatry, Vol. 17, 275-280,
45 1982), where mainly the destruction of cholinerg neurons (neurons which synthesise acetylcholine) can be observed in part because the increased production of free radicals. Another therapeutic benefit of lecithin is that it decreases the high cholesterol level of blood attenuating the risk of atherosclerosis (Lecithin and health care, Semmelweis Verlag, Hoya, Germany, 1985). Recently, the hepatoprotective effect of lecithin was discovered, which is
50 also used for therapeutic aim (Nutrition Review, Vol. 52, 327-339, 1994.).

In human beings the role of ascorbic acid otherwise vitamin-C was seen from its discovery (1928) to several decades in that it needs to the synthesis of collagen to prevention of scurvy. However, in recently more and more biological functions of ascorbic acid have become known mainly from that time when its antioxidant property and the presence of free
55 radicals in the living body were discovered.

Free radicals are chemically very active molecules because they have redundant electron. These molecules for example superoxide-anion (O_2^-), hydroxyl free radical (OH^\cdot) and free radical generating hydrogen-peroxide (H_2O_2) are formed during the normal metabolic processes of cells and can react with macromolecules (proteins, lipids and deoxyribonucleic
60 acid) changing hereby their structures. These changes may endanger the normal function of cell in great degree. For the sake of prevention of danger effect of free radicals cells have defensive mechanisms and protective agents such as ascorbic acid, which can adsorb and neutralise free electrons. The preventive effect of ascorbic acid is very important in stress situation, when the production of free radicals is augmented significantly because the

65 increased cell function. During the stress large volume of ascorbic acid is secreted in part with preventive aim from tissue stores to blood carrying it to different cells. However, if the degree of stress is higher than the possibilities of preventive mechanisms, for example stress is very high or chronic or the amount of ascorbic acid or other antioxidants are not enough, cells and tissues will be damage, which will manifest in the form of different diseases such as
70 atherosclerosis, decreased immune system activity, decreased viability of some neurons, Parkinson disease.

Ascorbic acid beside its free radical scavenger activity affects significantly the function of neurones, too, for example increases the secretion of acetylcholine and noradrenaline neurotransmitters as shown by in vitro neuronal study, and its large doses (1-3 g/day)
75 stimulate the effect of antipsychotic drugs in schizophrenic people (Progress in Neurobiology Vol. 43, 537-565, 1994).

Because ascorbic acid is essential for human organism and has positive effect in several diseases it is used alone and in combination with other active ingredients, for example with antipyretics (acetylsalicylic acid, paracetamol) and with metal ions (calcium, magnesium and
80 iron).

On the basis of biological knowledge of lecithin (L-alpha-phosphatidylcholine) and ascorbic acid it may be presumable that using them in combination they can complete each other's effects. However when we studied lecithin ascorbic acid combination in animal experiments we experienced surprising manner that mixing the two compounds in an
85 adequate ratio and giving it to rodents the effect of blend was more potent than the effect of components separately or the sum of the separated effect of components. This type of drug interaction is said to potentiating synergism, which was appeared in the increase of learning and physical performances of animals. The present invention therefore provides a drug combination, which comprises lecithin with 10.00-99.99 mass% L-alpha-phosphatidylcholine content and ascorbic acid in 1:1-30:1 mass ratio as active ingredients and one or more
90 pharmaceutically acceptable excipients, and provides a food product, a refreshing beverage or a concentrate for their production which comprises lecithin with 10.00-99.99 mass% L-alpha-phosphatidylcholine content and ascorbic acid in 1:1-30:1 mass ratio beside the known substances, and provides the use of these products the preservation of the physical and
95 cognitive performance of healthy people, the improvement of the physical and cognitive performance of healthy and sick people, the bracing of healthy and sick people and the increase of the physical and cognitive performance of sportsmen and sportswomen.

Studying the effect of combination and its components separately on the learning process of rats in water labyrinth (Brain Research Bulletin Vol. 45, 475-488, 1998) it was found that, rats treating with combination could find out from labyrinth with fewer error and in the first and third session in shorter time than vehicle (control) or only lecithin or only ascorbic acid treated animals (Table I.). Ascorbic acid and lecithin were ineffective when they were given separately (Table I.).

Table I. Effect of ascorbic acid (Vit.-C), lecithin and their combination on the learning process of normal rats in water labyrinth

	Number of session	Control	Vit.-C	Lecithin	Lecithin+Vit.-C
Dose (mg/kg/day) p.o.			2×10	2×30	2×(30+10)
Number of animals		10	10	10	10
Number of errors Mean±S.E.M.	1	16.0±0.39	16.4±0.45	15.8±0.29	13.2±0.46**†
	2	10.9±0.77	11.6±0.40	11.8±0.57	8.7±0.97†
	3	5.4±0.52	6.2±0.76	6.1±0.62	3.2±0.55*†
	4	2.9±0.72	3.9±0.35	3.2±0.59	2.6±0.67
Swimming time (sec) Mean±S.E.M.	1	138.4±8.95	140.3±9.54	129.5±7.61	110.7±8.87 ^{ab}
	2	90.7±7.80	89.9±6.22	80.1±5.54	75.9±8.59
	3	51.6±8.17	64.9±7.29	62.9±6.34	46.1±5.38 ^{ab}
	4	29.9±2.33	34.8±2.79	40.1±3.66 ^a	42.3±7.57

* p<0.05, ** p<0.01 compared with control by Mann-Whitney test

* p<0.05 compared with control by Student's t-test

† p<0.05 ‡ p<0.01 compared with Vit.-C and lecithin by Mann-Whitney test

* p<0.05 compared with Vit.-C by Student's t-test

^a p<0.05 compared with lecithin by Student's t-test

The composition of lecithin: 78.4% phospholipid (inside it 63.1% L-alpha-phosphatidylcholine, 24.1% L-alpha-phosphatidylethanolamine, 7.6% L-alpha-phosphatidilinositol, 4.6 % L-alpha-phosphatidic acid) and 20 % other lipid.

In water labyrinth learning test the combination prevented the increase of error number and swimming time caused by scopolamine a memory-destroying agent (Table II.). The two measured parameters were almost normalised by lecithin ascorbic acid blend (Table II.).

Studying the effect of combination and its components separately on the physical performance of mice in swimming test (Pharmacological Research, Vol. 23, 149-155, 1991) it was found that mice treating with the combination could swim longer time than vehicle (control) or only lecithin or only ascorbic acid treated animals (Table III.). In the study ascorbic acid was ineffective, when it was given in itself (Table III.).

Table II. Effect of lecithin ascorbic acid (Vit.-C) combination on the learning process of rats in scopolamine induced memory deficit in water labyrinth

	Number of session	Placebo	Scopolamine	Scopolamine
Dose (mg/kg/day) p.o.		control	control	Lecithin+Vit.-C 2×(30+10)
Number of animals		10	10	10
Number of errors Mean±S.E.M.	1	14.0±0.68	16.2±0.83	13.9±0.72
	2	6.6±0.76	11.6±0.52 ^{††}	8.8±0.59 ^{**}
	3	2.8±0.53	8.9±0.41 ^{††}	5.1±0.50 ^{**}
	4	2.0±0.33	6.3±0.26 ^{††}	3.9±0.48 ^{**}
Swimming time (sec) Mean±S.E.M.	1	96.4±8.44	114.7±9.45	78.1±5.31 ^{##}
	2	57.5±5.69	73.7±6.22	59.7±3.94
	3	37.6±4.08	66.7±6.47 ^{##}	41.7±3.21 ^{##}
	4	27.1±1.36	42.4±2.78 ^{##}	40.5±3.96

*p<0.05, **p<0.01 compared with scopolamine control by Mann-Whitney test

^{##}p<0.01 compared with scopolamine control by Student's t-test

^{††}p<0.01 compared with normal control by Mann-Whitney test

^{##}p<0.01 compared with normal control by Student's t-test

Table III. Effect of ascorbic acid (Vit.-C), lecithin and their combination on the swimming time of normal mice

			Number of days				
	Dose mg/kg/day p.o.	Number of animals	1. day starting values	2. day	3. day	4. day	5. day
			Swimming time (sec) mean \pm S.E.M.				
Control		10	77.4 \pm 4.5	80.4 \pm 3.9	82.2 \pm 5.9	84.6 \pm 7.3	84.0 \pm 6.8
Vit.-C	2 \times 10	10	76.2 \pm 3.7	70.8 \pm 3.8	82.2 \pm 3.4	88.8 \pm 6.1	90.0 \pm 6.9
Lecithin	2 \times 30	10	70.2 \pm 3.6	100.2 \pm 6.9*	20.6 \pm 7.8**	126.0 \pm 8.0**	126.6 \pm 9.1**
Lecithin+Vit.-C	2 \times (30+10)	10	72.0 \pm 3.2	106.8 \pm 11.9*	32.6 \pm 9.4**	144.0 \pm 13.9**	141.0 \pm 13.0**

* p<0.05, ** p<0.01 compared with control by Student's t-test

On the basis of the results of these experiments it can be established that lecithin is mixed with ascorbic acid in an appropriate ratio and using this blend in suitable formula we can gain such a drug product, which can be applied more favourable and more effectively to those areas where these agents are used alone or maybe together but no optimised ratio. In addition, in the combination the smaller dose of lecithin and ascorbic acid can produce as effect as ingredients separately decreasing by this means the side effects of ingredients. The

higher dose of ascorbic acid may promote kidney stone while the high dose of lecithin, higher than 25 g/day may produce loss of appetite, nausea, stomach puffing and diarrhoea.

Beside the general tonic action of lecithin ascorbic acid combination (increase of physical and cognitive performance) it can use in the protection of heart and vascular system and in their diseases or in the protection of brain functions and in their diseases or in the protection of liver and in its diseases or in vitamin-C or in acetylcholine deficiency or in the prevention of bacterial or virus infections and in their diseases or in the prevention of diabetes complications and in this disease or in the stimulating of immune system.

In the combination the mass ratio of lecithin and ascorbic acid can change from 1:1 to 30:1, while the daily doses of components depending on the aim of application can vary between 1.5-500 mg/kg in the case of lecithin and between 1-60 mg/kg in the case of ascorbic acid dividing these doses into suitable portion.

In a preferable drug formula lecithin is present in well dispersed solid or emulsified or liposome or dissolved form, which contain ascorbic acid solid or dissolved form. The blend of two ingredients can be used in oral and parenteral drug formula according to need together with auxiliary materials, which are used for drug formulation. In these drug forms the mass of two ingredients can vary as a dose between 1-1000 mg respectively.

In what follows mentioned according to Example 1 produced capsule can be favourable used for prevention of illnesses or physical and cognitive exhaustion in 2×2 or 3×2 amount in healthy people. In illnesses as an additive treatment for tonic aim 2×2, 3×2 or 4×2 capsule and for the sake of tendentious effect, for example for the prevention of diabetes caused lipidperoxidation or ageing memory disorders or Alzheimer disease or Parkinson diseases 3×2, 4×2 or 4×3 capsule may be favourable. In some liver diseases where the lower ascorbic acid dose is favourable 4×3 or 4×4 capsule can be therapeutic worth from that product, which is made according to example 2 with 20:1 mass ratio of lecithin ascorbic acid. If the oral dosing can not be solved, a treatment of injections can come to the front.

The following non-limiting examples further illustrate the invention:

Example 1

Formulation of lecithin and ascorbic acid combination in 3:1 ratio in hard gelatine capsule

2 part by mass lecithin is suspended into 4.5 part by mass sunflower oil at 45-50 °C. Sunflower oil has to contain as an antioxidant 0.065 part by mass vitamin E oil. After the suspending of lecithin the oil is cooled on 20 °C and 0.66 part by mass well pulverised

(micronized) ascorbic acid is suspended into it. Finally this oil is formulated in 0.7 ml hard gelatine capsules.

Example 2

Formulation of lecithin and ascorbic acid combination in 20:1 ratio in hard gelatine capsule

2 part by mass lecithin is suspended into 4.5 part by mass sunflower oil at 45-50 °C. Sunflower oil has to contain as an antioxidant 0.065 part by mass vitamin E oil. After the suspending of lecithin the oil is cooled on 20 °C and 0.1 part by mass well pulverised (micronized) ascorbic acid is suspended into it. Finally this oil is formulated in 0.7 ml hard gelatine capsules.

Example 3

Formulation of lecithin and ascorbic acid combination in oil injection

Among sterile circumstance using sterile ingredients without pyrogen the injection is made according to the example 1 and the oil suspension of the combination is filled into 2 ml ampoule. This injection can be applied subcutan or intramuscular route.

Example 4

Formulation of lecithin and ascorbic acid combination in oil/water type injection

Among sterile circumstances using sterile ingredients without pyrogen 3 part by mass lecithin is suspended into 4.5 part by mass sunflower oil at 45-50 °C. Sunflower oil has to contain as an antioxidant 0.075 part by mass vitamin E oil. After the suspending of lecithin the oil is cooled on 20 °C, and it is emulsified in distilled water, which contains 0.66 part by mass ascorbic acid. The water phase is 1.5 times of oil phase. In this emulsion the emulsifier is itself lecithin. The emulsion is filled into 2 ml ampoule and can be applied subcutan or intramuscular route.

CLAIMS

1. A drug combination which comprises lecithin with 10.00-99.99 mass% L-alpha-phosphatidylcholine content and ascorbic acid in 1:1-30:1 mass ratio as active ingredients and one or more pharmaceutically acceptable excipients.
- 5 2. A food product, a refreshing beverage or a concentrate for their production which comprises lecithin with 10.00-99.99 mass% L-alpha-phosphatidylcholine content and ascorbic acid in 1:1-30:1 mass ratio beside the known substances.
3. Use of drug combination according to Claim 1 wherein the preservation of the physical and cognitive performance of healthy people, the improvement of the physical and cognitive performance of healthy and sick people, the bracing of healthy and sick
10 people and the increase of the physical and cognitive performance of sportsmen and sportswomen.
4. Use of drug combination according to Claim 1 wherein the replacement of acetylcholine or vitamin C deficiency, the prevention or curing of diseases connecting with
15 lipidperoxidation as for example Alzheimer disease, Parkinson disease, constriction of the brain, heart and periphery arteries, atherosclerosis caused diseases, high blood lipid and cholesterol concentration, bacterial and viral infections, liver diseases and diabetes.
5. Use of drug combination according to Claim 1 wherein oral (tablet, soft or hard gelatine capsule etc.) and parenteral (injection, sublingual tablet, ointment, suppository, plaster
20 etc.) drug forms.
6. Use of drug combination according to Claim 1 wherein jointly use with any itself known ingredient or vitamin in one unit.